WECARE ANTI-DANDRUFF- antidandruff shampoo liquid
Dynarex Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

1398 Antidandruff Shampoo NDC 67777-139-80

Active ingredient

Pyrithione Zinc 0.5%

Purpose

Anti-Dandruff

Use

Helps eliminate irritation and flaking associated with dandruff

Warnings

For External Use Only

When using this product

- Avoid contact with eyes
- If contact occurs, rinse eyes with plenty of water

Stop use and ask a doctor if

condition does not improve or worsens after regular use of this product as directed.

If swallowed, get medical help or contact a Poison Control Center immediately.

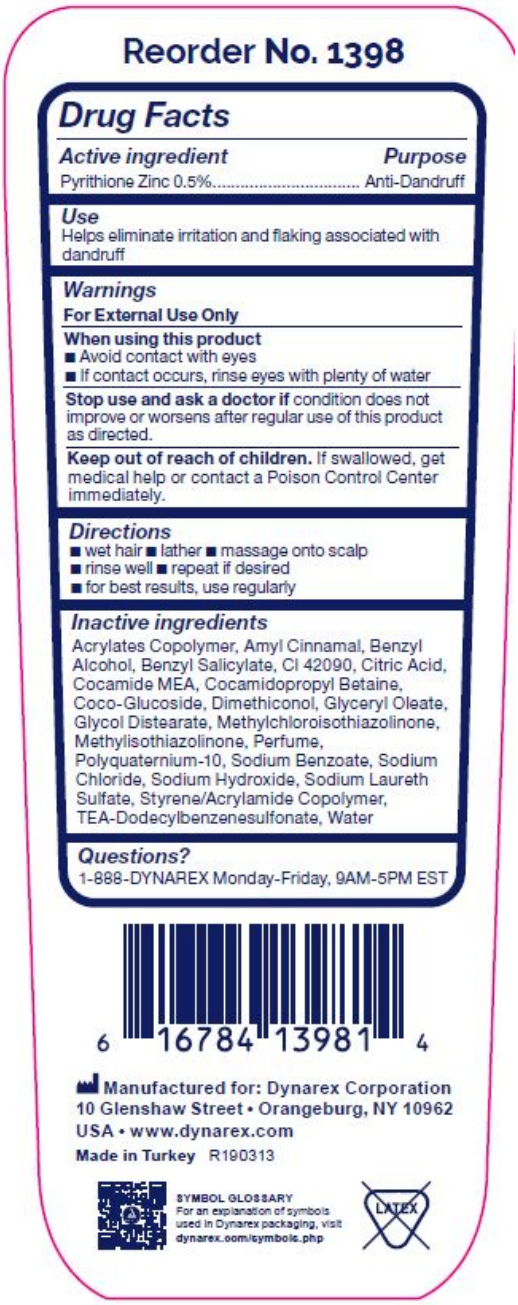
Directions

- wet hair
- lather
- massage onto scalp
- rinse well
- repeat if desired
- for best results, use regularly

Inactive ingredients

Water, Sodium Laureth Sulfate, Cocamide MEA, Cocamidopropyl Betaine, Sodium Chloride, Glycol Distearate, Acrylates Copolymer, Coco-Glucoside, Styrene/Acrylates Copolymer, Dimethiconol, Glyceryl Oleate, Polyquaternium-10, Citric Acid, TEA-dodecyl-benzene-sulfonate, Sodium Hydroxide, Benzyl Alcohol, Sodium Benzoate, Methylchloroisothiazolinone, Methylisothiazolinone, Amyl Cinnamal, Benzyl Salicylate, CI42090

1398 Label



WECARE ANTI-DANDRUFF

antidandruff shampoo liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-139
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	5 g in 1000 g

Inactive Ingredients	
Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
.ALPHA.-AMYLCINNAMALDEHYDE (UNII: WC51CA3418)	
WATER (UNII: 059QF0KO0R)	
STYRENE/ACRYLAMIDE COPOLYMER (MW 500000) (UNII: 5Z4DPO246A)	
DIMETHICONOL (2000 CST) (UNII: T74O12AN6Y)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
GLYCERYL MONOOLEATE (UNII: C4YAD5F5G6)	
POLYQUATERNIUM-10 (1000 MPA.S AT 2%) (UNII: GMR4PEN8PK)	
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO GLUCOSIDE (UNII: ICS790225B)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-139-80	12 in 1 CASE	10/11/2018	
1		335 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	10/11/2018	

Labeler - Dynarex Corporation (008124539)

Revised: 6/2019 Dynarex Corporation